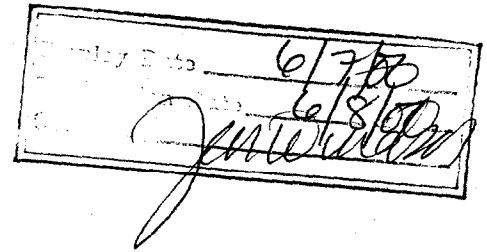


DMB



**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**Technical Electronic Product Radiation Safety Standards Advisory Committee;  
Notice of Meeting**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

---

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

*Name of Committee:* Technical Electronic Product Radiation Safety Standards Advisory Committee.

*General Function of the Committee:* To provide advice on technical feasibility, reasonableness, and practicality of performance standards for electronic products to control the emission of radiation under 42 U.S.C. 263f(f)(1)(A).

*Date and Time:* The meeting will be held on June 21, 2000, 8:30 a.m. to 5 p.m., and June 22, 2000, 8:30 a.m. to 12 noon.

*Location:* Quality Suites, Potomac II and III, 3 Research Ct., Rockville, MD.

*Contact Person:* Orhan H. Suleiman, Center for Devices and Radiological Health (HFZ-240), Food and Drug Administration, 1350 Piccard Dr., Rockville, MD 20850, 301-594-3332, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12399. Please call the Information Line for up-to-date information on this meeting.

*Agenda:* On June 21, 2000, the committee will discuss: (1) A reproposal for amendments to the performance standard for lasers (part 1040 (21 CFR part 1040)), (2) amendments being considered for the sunlamps standard (part 1040), and hear (3) a presentation addressing nonmedical ionizing radiation security systems, (4) a presentation concerning computed tomography

fluoroscopy (CT) and the Year 2000 Nationwide Evaluation of X-Ray Trends Survey of CT. On June 22, 2000, the committee will hear: (1) An update on the reengineering of the radiological health program at the Center for Devices and Radiological Health, (2) a presentation on manufacturers' requirements and the medical device approval process, and (3) a presentation regarding how ultrasound diathermy (21 CFR part 1050) is regulated by FDA.

*Procedure:* Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by June 16, 2000. Oral presentations from the public will be scheduled on June 21, 2000, between approximately 11 a.m. and 11:20 a.m., and between approximately 1:45 p.m. and 2:05 p.m. Oral presentations from the public will be scheduled on June 22, 2000, between approximately 10:30 a.m. and 11 a.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before June 16, 2000, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

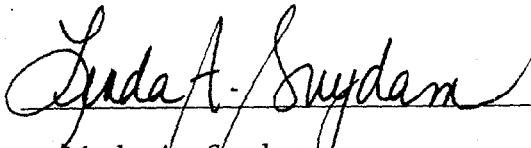
FDA regrets that it was unable to publish this notice 15 days prior to the June 21, 2000, Technical Electronic Product Radiation Safety Standards Advisory Committee meeting. Because the agency believes there is some urgency to bring these issues to public discussion and qualified members of the Technical Electronic Product Radiation Safety Standards Advisory Committee were available at this time, the Commissioner of Food and Drugs concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app.

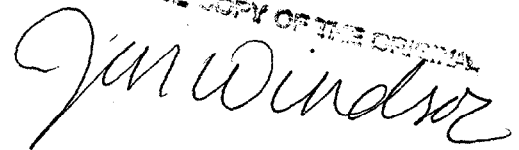
2).

Dated: 6/1/00

June 1, 2000

  
Linda A. Suydam  
Senior Associate Commissioner

CERTIFIED TO BE A TRUE COPY OF THE ORIGINAL



[FR Doc. 00-???? Filed ??-??-00; 8:45 am]

BILLING CODE 4160-01-F